

(19)



Europäisches Patentamt
European Patent Office
Office européen des brevets



(11)

EP 1 031 579 A2

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication:
30.08.2000 Bulletin 2000/35

(51) Int Cl.7: **C08F 2/50, G03F 7/029,
G03F 7/00**

(21) Application number: **00103990.8**

(22) Date of filing: **25.02.2000**

(84) Designated Contracting States:
**AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE**
Designated Extension States:
AL LT LV MK RO SI

(30) Priority: **26.02.1999 JP 4974299
04.05.1999 US 132449 P**

(71) Applicant: **SHOWA DENKO KABUSHIKI KAISHA
Minato-ku, Tokyo (JP)**

(72) Inventors:
• Katoh, Tsuyoshi, c/o Showa Denko K.K.
Cent.Res.Lab
Kawasaki-shi, Kanagawa-ken (JP)
• Ogata, Tomonari, c/o Showa Denko K.K.
Cent.Res.Lab
Kawasaki-shi, Kanagawa-ken (JP)

(74) Representative: **Strehl Schübel-Hopf & Partner
Maximilianstrasse 54
80538 München (DE)**

(54) **Photopolymerization initiator for color filter, photosensitive coloring composition, and color filter**

(57) A photopolymerization initiator for producing a color filter including a sensitizer (A) which absorbs light having a wavelength of 300 to 500 nm, an organic boron complex (B), and a thiol-group-containing compound (C); a photosensitive coloring composition including a transparent polymer (D) which is soluble to a solvent or

an aqueous alkaline solution, a compound having ethylenic unsaturated bonds (E), a pigment (F), and the photopolymerization initiator; a color filter having a pixel formed by use of the coloring composition on a transparent substrate; and a method for producing the color filter.

EP 1 031 579 A2

Description**FIELD OF THE INVENTION**

5 [0001] The present invention relates to a high-sensitivity photopolymerization initiator for producing a color filter; to a high-quality photosensitive coloring composition for producing a color filter, which can be developed with a solvent or an aqueous alkaline solution after irradiation with UV light; to a color filter which is used in devices such as a solid state pickup device and a liquid crystal display device; and to a method for producing the color filter.

10 BACKGROUND OF THE INVENTION

[0002] In recent years, demand has arisen for a color filter used in a device such as a liquid crystal display device, where the filter is thin and defect-free; has properties such as heat and light resistance; and has optical properties such as high transparency and high contrast in accordance with the trend toward reducing the thickness and increasing 15 the width of the device. In addition, increase of the yield of a production step and reduction in the number of production steps are more and more required, along with a demand for reducing production cost of a color filter. A color filter per se comprises a transparent substrate such as a glass plate on which fine stripes of two or more hues are disposed in a parallel or crossing manner or fine pixels are disposed in lateral and longitudinal directions in a predetermined pattern. Each pixel has a fine shape and a dimension of some 10 µm to some 100 µm, and the pixels for each hue must be 20 arrayed according to a predetermined order. Thus, a variety of methods for producing a color filter have been proposed. Of these, from many viewpoints, such as the precision of the produced filter and production cost, a pigment-dispersion method employing a photosensitive coloring composition in which a pigment is dispersed has gained wide acceptance.

[0003] The pigment-dispersion method comprises applying, on a substrate, a photosensitive coloring composition containing a pigment, a resin serving as a matrix, and a cross-linking agent; and forming each coloring layer in a 25 predetermined pattern through photolithography. More specifically, a color filter is produced through the following steps: applying the aforementioned photosensitive coloring composition for achieving one filter color on a transparent substrate such as a glass plate; exposing patternwise; developing the unexposed portion with a solvent or an aqueous alkaline solution to thereby form a pattern of a first color; and sequentially repeating the procedure for another filter color. A negative-type photosensitive coloring composition is predominantly employed as the above described coloring 30 composition, in view of properties of the produced color filter and a wide range of usable materials. Currently, during development, a solvent is used very seldom, from the viewpoints of environmental problems; instead, an aqueous alkaline solution is predominantly used. Exposure is typically carried out by use of UV light. In the above production steps, only a predetermined portion is irradiated with light such as UV light, to thereby cause a chemical reaction; i.e., radical polymerization, exclusively in the exposed portion.

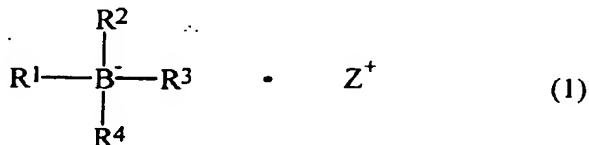
[0004] Since a negative-type photosensitive composition predominantly undergoes radical polymerization, the sensitivity of the composition may be reduced through inhibition of polymerization caused by oxygen in the air. In order to solve this problem, for example, the applied photosensitive composition is irradiated with light after an oxygen-shielding film such as a poly(vinyl alcohol) film is formed on the composition. However, recently, demand has arisen for a photopolymerization initiator for producing a color filter, where the initiator has such high sensitivity that additional treatment 40 such as the provision of an oxygen-shielding film can be eliminated.

[0005] In this connection, Japanese Unexamined Patent Application, First Publication, No. 6-201913 discloses a photopolymerization initiator comprising solely a triazine compound, and a photopolymerization initiator comprising a triazine compound and an imidazole compound in combination. Japanese Unexamined Patent Application, First Publication, No. 10-253813 discloses a photopolymerization initiator comprising a polyfunctional thiol compound and at 45 least one compound selected from a biimidazole compound, a titanocene compound, a triazine compound, and an oxadiazole compound. Photopolymerizable compositions containing these photopolymerization initiators for achieving red, green, and blue colors have a certain degree of sensitivity. However, a photopolymerizable composition for achieving a black color, particularly a photopolymerizable composition containing a black pigment, has drawbacks, such as a considerable reduction in sensitivity due to a shielding effect of the black pigment during exposure, increase of 50 exposure time, and insufficient curing, even when the composition contains the above-described polymerization initiator.

SUMMARY OF THE INVENTION

55 [0006] The present inventors have found that a photopolymerization initiator comprising a sensitizer (A) which absorbs light having a wavelength of 300-500 nm, an organic boron complex (B), and a thiol-group-containing compound (C) is suitable for producing a color filter, and that the above problems are solved by use of the initiator. The present invention has been accomplished on the basis of this finding.

- [0007] In view of the foregoing, an object of the present invention is to provide a high-sensitivity photopolymerization initiator for producing a color filter.
- [0008] Another object of the present invention is to provide a photosensitive coloring composition containing the initiator.
- 5 [0009] Another object of the present invention is to provide a color filter exhibiting excellent resistance to a variety of phenomena.
- [0010] Another object of the present invention is to provide a method for producing the color filter.
- [0011] Accordingly, the present invention provides a photopolymerization initiator for producing a color filter comprising a sensitizer (A) which absorbs light having a wavelength of 300 to 500 nm; an organic boron complex (B) represented by formula (1):
- 10



15 wherein each of R¹, R², R³, and R⁴ represents an alkyl group, an aryl group, an aralkyl group, an alkenyl group, a heterocyclic group, or an alicyclic group; and Z⁺ represents an ammonium cation, a sulfonium cation, an oxosulfonium cation, a pyridinium cation, a phosphonium cation, an oxonium cation, or an iodonium cation; and a thiol-group-containing compound (C).

20 [0012] Preferably, the thiol-group-containing compound (C) is at least one species selected from 2-mercaptopbenzothiazole, trimethylolpropane tris(2-mercaptopacetate), trimethylolpropane tris(3-mercaptopropionate), pentaerythritol tetrakis(2-mercaptopacetate), and pentaerythritol tetrakis(3-mercaptopropionate).

25 [0013] Preferably, the sensitizer (A) contains a benzophenone compound and/or a biimidazole compound.

[0014] The present invention also provides a photosensitive coloring composition for producing a color filter, which comprises a transparent polymer (D) soluble in a solvent or an aqueous alkaline solution, a compound having an ethylenic unsaturated bond (E), a pigment (F), and the above-described photopolymerization initiator.

30 [0015] The present invention also provides a color filter having pixels formed on a transparent substrate by use of the photosensitive coloring composition.

[0016] The present invention also provides a method for producing a color filter comprising forming a layer of the photosensitive coloring composition on a transparent substrate and photocuring exposed portions of the layer of the composition through exposure.

35

DESCRIPTION OF THE PREFERRED EMBODIMENTS

[0017] The present invention will next be described in more detail. When R¹, R², R³, and R⁴ in formula (1) represent alkyl groups, and the alkyl groups may have substituents. Specifically, a C₁-C₁₂ linear or branched alkyl group which may have a substituent is preferred. Examples of suitable alkyl groups include a methyl group, an ethyl group, an n-propyl group, an isopropyl group, an n-butyl group, an isobutyl group, a sec-butyl group, a tert-butyl group, an n-pentyl group, an n-hexyl group, an n-octyl group, a dodecyl group, a cyanomethyl group, a 4-chlorobutyl group, a 2-diethylaminoethyl group, and a 2-methoxyethyl group.

[0018] When R¹, R², R³, and R⁴ in formula (1) represent aryl groups, the aryl groups may have substituents. Specifically, the groups are aryl groups which may be substituted. Examples of suitable aryl groups include a phenyl group, a tolyl group, a xylyl group, a mesityl group, a 4-methoxyphenyl group, a 2-methoxyphenyl group, a 4-n-butylphenyl group, a 4-tert-butylphenyl group, a naphthyl group, a 4-methylnaphthyl group, a 4-ethylnaphthyl group, an anthryl group, a phenanthryl group, a 4-nitrophenyl group, a 4-trifluoromethylphenyl group, a 4-fluorophenyl group, a 4-chlorophenyl group, and a 4-dimethylaminophenyl group.

[0019] When R¹, R², R³, and R⁴ in formula (1) represent aralkyl groups, the aralkyl groups may have substituents. Specifically, the groups are aralkyl groups which may be substituted. Examples of suitable aralkyl groups include a benzyl group, a phenethyl group, a 1-naphthylmethyl group, a 2-naphthylmethyl group, and a 4-methoxybenzyl group.

[0020] When R¹, R², R³, and R⁴ in formula (1) represent alkenyl groups, the alkenyl groups may have substituents. Specifically, the groups are alkenyl groups which may be substituted. Examples of suitable alkenyl groups include a vinyl group, a propenyl group, a butenyl group, and an octenyl group.

[0021] When R¹, R², R³, and R⁴ in formula (1) represent heterocyclic groups, the heterocyclic groups may have substituents. Specifically, the groups are heterocyclic groups which may be substituted. Examples of such heterocyclic groups include a pyridyl group, a 4-methylpyridyl group, a quinolyl group, and an indolyl group.

- [0022] When R¹, R², R³, and R⁴ in formula (1) represent alicyclic groups, the alicyclic groups may have substituents. Specifically, the groups are alicyclic groups which may be substituted. Examples of such alicyclic groups include a cyclohexyl group, a 4-methylcyclohexyl group, a cyclopentyl group, and a cycloheptyl group.
- [0023] When Z⁺ in formula (1) represents an ammonium cation, examples thereof include a tetramethylammonium cation, a tetraethylammonium cation, a tetra-n-propylammonium cation, a tetra-n-butylammonium cation, an n-butyltriphennylammonium cation, a tetraphenylammonium cation, and a benzyltriphenylammonium cation.
- [0024] When Z⁺ in formula (1) represents a sulfonium cation, examples thereof include a triphenylsulfonium cation, a tri(4-tolyl)sulfonium cation, and a 4-tert-butylphenyldiphenylsulfonium cation.
- [0025] When Z⁺ in formula (1) represents an oxosulfonium cation, examples thereof include a triphenyloxosulfonium cation, a tri(4-tolyl)oxosulfonium cation, and a 4-tert-butylphenyldiphenyloxosulfonium cation.
- [0026] When Z⁺ in formula (1) represents an oxonium cation, examples thereof include a triphenyloxonium cation, a tri(4-tolyl)oxonium cation, and a 4-tert-butylphenyldiphenyloxonium cation.
- [0027] When Z⁺ in formula (1) represents a pyridinium cation, examples thereof include an N-methylpyridinium cation and an N-n-butylpyridinium cation.
- [0028] When Z⁺ in formula (1) represents a phosphonium cation, examples thereof include a tetramethylphosphonium cation, a tetra-n-butylphosphonium cation, a tetra-n-octylphosphonium cation, a tetraphenylphosphonium cation, and a benzyltriphenylphosphonium cation.
- [0029] When Z⁺ in formula (1) represents an iodonium cation, examples thereof include a diphenyliodonium cation, a di(4-methylphenyl)iodonium cation, and a di(4-tert-butylphenyl)iodonium cation.
- [0030] Examples of organic boron complexes represented by formula (1) include tetramethylammonium n-butyltriphennyborate, tetraethylammonium isobutyltriphennyborate, tetra-n-butylammonium methyltriphennyborate, tetra-n-butyltriphennyborate, tetra-n-butylammonium n-octyl-triphennyborate, tetra-n-butylammonium n-butyltriphennyborate, tetra-n-butylammonium n-butyltrinaphthylborate, tetra-n-butylammonium n-butyltrinaphthylborate, tetra-n-butylammonium n-octyltrinaphthylborate, tetra-n-butylammonium methyltri(4-methylnaphthyl)borate, tetra-n-butylammonium n-butyltri(4-methylnaphthyl)borate, triphenylsulfonium n-butyltriphennyborate, triphenylsulfonium n-butyltrinaphthylborate, triphenyloxosulfonium n-butyltriphennyborate, triphenyloxosulfonium n-butyltrinaphthylborate, triphenyloxonium n-butyltriphennyborate, N-methylpyridinium n-butyltriphennyborate, tetraphenylphosphonium n-butyltriphennyborate, and diphenyliodonium n-butyltriphennyborate.
- [0031] In general, the organic boron complex represented by formula (1) absorbs substantially no light having a wavelength of 300 nm or longer. Thus, when the complex is used alone, it is not sensitive to light from a typical UV lamp. However, the combination of the complex and a sensitizer has remarkably high sensitivity.
- [0032] In the present invention, the organic boron complex (B) is incorporated typically in an amount of 1 to 60 % by weight based on the compound having ethylenic unsaturated bonds, preferably 5 to 30 % by weight. When the amount of the organic boron complex (B) is insufficient, polymerization proceeds insufficiently, whereas when the amount of the organic boron complex (B) is excessive, the stability of the composition may decrease, to thereby cause an economically disadvantageous situation.
- [0033] Examples of the sensitizer (A) which absorbs light having a wavelength of 300 to 500 nm used in the invention include a photopolymerization initiator, a sensitizer, and a dye absorbing light having a wavelength in the range. Examples of the sensitizers include cationic dyes; benzophenones; acetophenones; benzoins; thioxanthones; imidazoles; biimidazoles; coumarins; ketocoumarins; triphenylpyryliums; triazines; benzoic acids; and maleimides. In addition, acylphosphine oxide, methyl phenyl glyoxylate, an α -acyloxime ester, benzil, camphorquinone, and ethylanthraquinone may also be employed. These compounds may be used alone or as a combination of two or more thereof.
- [0034] Specific examples of the sensitizers include cationic dyes such as Basic Yellow 1, 11, 13, 21, 28, 29, 36, 51 and 63, Basic Orange 21 and 22, benzophenone, 4-methylbenzophenone, 4-fluorobenzophenone, 4-chlorobenzophenone, 4-bromobenzophenone, 4-morpholinobenzophenone, 4-dimethylaminobenzophenone, 4,4'-bis(dimethylamino)benzophenone, 4,4'-bis(diethylamino)benzophenone, 4-dimethylaminoacetophenone, 4-diethylaminoacetophenone, 2-hydroxy-2-phenylacetophenone, 2,4-diethylthioxanthone, 2-methylthioxanthone, N-methylimidazole, N-phenylimidazole, 2,2'-bis(2-chlorophenyl)-4,4',5,5'-tetraphenyl-1,2'-biimidazole (hereinafter referred to as HABI), 2,2'-bis(2-chlorophenyl)-4,4',5,5'-tetra(ethoxyphenyl)-1,2'-biimidazole, 2,2'-bis(2-chlorophenyl)-4,4',5,5'-tetra(4-bromophenyl)-1,2'-biimidazole, 2,2'-bis(2,4-dichlorophenyl)-4,4',5,5'-tetraphenyl-1,2'-biimidazole, 2,2'-bis(2-chlorophenyl)-4,4',5,5'-tetra(3-methoxyphenyl)-1,2'-biimidazole, 2,2'-bis(2-methylphenyl)-4,4',5,5'-tetraphenyl-1,2'-biimidazole, coumarin, 7-diethylaminocoumarin, ethyl 4-dimethylaminobenzoate, N-methylmaleimide, and N-phenylmaleimide. These compounds may be used alone or as a combination of two or more thereof. Considering stability of the initiator and a composition and ability to initiate polymerization, benzophenone compounds, benzoic acids, and biimidazole compounds are preferably used.
- [0035] In the present invention, the sensitizer (A) which absorbs light having a wavelength of 300 to 500 nm is incorporated typically in an amount of 1 to 60 wt.% based on the compound having ethylenic unsaturated bonds,

preferably 2 to 30 wt.%. When the amount of the sensitizer (A) is insufficient, the sensitizing effect may not be obtained, whereas when the amount of the sensitizer (A) is excessive, the sensitizer (A) absorbs light and thus light transmittance efficiency may become poor, and polymerization initiation efficiency may decrease, which is unsatisfactory.

[0036] The thiol-group-containing compound (C) used in the present invention is a thiol-group-containing compound in the molecule. Examples of suitable thiol compounds include 2-mercaptopbenzothiazole, 2-mercaptopbenzimidazole, 2-mercaptopbenzoxazole, 5-chloro-2-mercaptopbenzothiazole, 2-mercaptop-5-methoxybenzothiazole, 5-methyl-1,3,4-thiadiazole-2-thiol, 5-mercaptop-1-methyltetrazole, 3-mercaptop-4-methyl-4H-1,2,4-triazole, 2-mercaptop-1-methyl-imidazole, 2-mercaptopthiazoline, octanethiol, hexanedithiol, decanedithiol, 1,4-dimethylmercaptopbenzene, butanediol bis(2-mercaptopacetate), butanediol bis(3-mercaptopropionate), ethylene glycol bis(2-mercaptopacetate), ethylene glycol bis(3-mercaptopropionate), trimethylolpropane tris(2-mercaptopacetate), trimethylolpropane tris(3-mercaptopropionate), pentaerythritol tetrakis(2-mercaptopacetate), pentaerythritol tetrakis(3-mercaptopropionate), and trishydroxyethyl tris(3-mercaptopropionate). In considering polymerization initiation ability, particularly, 2-mercaptopbenzothiazole, 2-mercaptopbenzimidazole, 2-mercaptopbenzoxazole, trimethylolpropane tris(2-mercaptopacetate), trimethylolpropane tris(3-mercaptopropionate), pentaerythritol tetrakis(2-mercaptopacetate), and pentaerythritol tetrakis(3-mercaptopropionate) are preferred. These compounds having a thiol group may be used alone or as a combination of two or more thereof.

[0037] In the present invention, the thiol-group-containing compound (C) is incorporated typically in an amount of 1 to 60 wt.% based on the compound having ethylenic unsaturated bonds, preferably 5 to 30 wt.%. When the amount of the thiol-group-containing compound (C) is insufficient, polymerization may not be effectively initiated, whereas when the amount of the thiol-group-containing compound (C) is excessive, the initiation performance may not increase and the addition disadvantageously affects the properties of the cured product.

[0038] In the present invention, the combination of the sensitizer (A) which absorbs light having a wavelength of 300 to 500 nm, the organic boron complex (B) represented by formula (1), and the thiol-group-containing compound (C) enhances polymerization performance.

[0039] In the above combination, preferably, the organic boron complex (B) is used in an amount from 5 to 500% by weight, based on the sensitizer (A) and the thiol-group-containing compound (C) is used in an amount from 10 to 500% by weight, based on the organic boron complex (B). Furthermore, more preferably, the organic boron complex (B) is used from 20 to 300% by weight, based on the sensitizer (A) and the thiol-group-containing compound (C) is used from 50 to 200% by weight, based on the organic boron complex (B). When the amount of the organic boron complex (B) is insufficient, the curing of the composition is insufficient, whereas when the amount of the organic boron complex (B) is excessive, the storing stability of the compound may decrease. Furthermore, when the amount of the thiol-group-containing compound (C) is insufficient, the curing of the composition is insufficient, whereas when the amount of the thiol-group-containing compound (C) is excessive, properties such as the water resistance and the mechanical properties, may decrease.

[0040] The transparent polymer (D) which is soluble in a solvent or an aqueous alkaline solution used in the present invention is a polymer with high transparency. Accordingly, a film formed of the polymer having a thickness of 1 μm preferably has a percent transmission of 80% or more, more preferably 95% or more within a visible range of 400 to 700 nm, and the polymer can be dissolved in a developer (a solvent or an aqueous alkaline solution). Examples of the polymer include thermosetting resins, thermoplastic resins, and photosensitive resins. These polymers may be used alone or as a combination of two or more thereof. In a post-treatment process for producing a color filter, the polymer is treated at high temperature or with a variety of solvents and chemicals, and therefore, the transparent polymer (D) having excellent heat resistance and resistance to solvents and chemicals is preferably used.

[0041] The compound (E) having ethylenic unsaturated bonds used in the present invention may be a compound which can undergo radical polymerization (or cross-linking) and is typically called a monomer or oligomer. Examples of suitable compounds include (meth)acrylic acid; (meth)acrylic acid esters such as methyl (meth)acrylate, butyl (meth)acrylate, benzyl (meth)acrylate, 2-hydroxyethyl (meth)acrylate, and 2-hydroxypropyl (meth)acrylate; ethylene glycol di(meth)acrylate; pentaerythritol tri(meth)acrylate; styrene; divinylbenzene; (meth)acrylamide; vinyl acetate; N-hydroxymethyl(meth)acrylamide; dipentaerythritol hexaacrylate; melamine acrylate; and an epoxy acrylate pre-polymer. Considering exposure sensitivity and resistance to a variety of phenomena after curing, multifunctional (meth)acrylic monomers are preferably used.

[0042] As used herein, the term "(meth)acryl" refers to both "methacryl" and "acryl."

[0043] Widely used hues of color filters include additive color mixing types such as red, green, and blue; subtractive color mixing types such as cyan, magenta, and yellow; and black used in a black matrix portion. Coloring agents include dyes and pigments, but pigments are used in the present invention considering heat-resistance and photo-resistance as described above. In order to obtain suitable spectra, two or more pigments are used in combination. For example, a blue spectrum is obtained from the combination of a cyan pigment and a violet pigment; a green spectrum is obtained from the combination of a blue pigment and a yellow pigment; and a red spectrum is obtained from the combination of a red pigment and a yellow or an orange pigment.

[0044] Examples of the pigment (F) used in the present invention, represented by color index numbers, include C. I. Pigment Yellow 12, 13, 14, 17, 20, 24, 55, 83, 86, 93, 109, 110, 117, 125, 137, 139, 147, 148, 153, 154, 166, 168, 217, 220, 223, 224, 226, 227, 228, 240, C. I. Pigment Violet 19, 23, 29, 30, 37, 40, 50, C. I. Pigment Blue 15, 15:1, 15:4, 15:6, 22, 60, 64, C. I. Pigment Green 7, 36, C. I. Pigment Brown 23, 25, 26, C. I. Pigment Black 7, and Titanium Black.

[0045] In the photosensitive coloring composition of the present invention, the pigment should be sufficiently dispersed. In addition, the photosensitive coloring composition should be applied to a transparent substrate to a thickness of 1 to 3 µm. Therefore, in order to impart suitable applicability to the composition, in general, the viscosity of the composition is controlled by use of a solvent. Examples of suitable solvents include methanol, ethanol, toluene, cyclohexane, isophorone, cellosolve acetate, diethylene glycol dimethyl ether, ethylene glycol diethyl ether, xylene, ethylbenzene, methyl cellosolve, ethyl cellosolve, butyl cellosolve, propylene glycol monomethyl ether, isoamyl acetate, ethyl lactate, methyl ethyl ketone, acetone, and cyclohexanone. These solvents may be used alone or as a combination of two or more thereof.

[0046] The photosensitive coloring composition is produced by use of any of a variety of means for dispersion, such as a three-roll mill, a two-roll mill, a sand mill, an attrition mill, a ball mill, a kneader, and a paint shaker. In order to prevent gelation attributed to polymerization during the dispersion process, a polymerization inhibitor may be added, and a monomer and the photopolymerization initiator may be added after dispersion of the pigment. Furthermore, in order to facilitate the dispersion of the pigment, an appropriate dispersing aid may be added. Since a dispersing aid facilitates dispersion and prevents re-aggregation of the pigment, a color filter of excellent transparency can be produced.

[0047] In general, the color filter of the present invention is produced by applying the photosensitive coloring composition of the present invention to a transparent substrate such as a glass substrate, and photocuring the composition through exposure. In general, the color filter is produced by the following steps:

in a first step, a photosensitive color resin layer is formed on a transparent substrate;
 in a second step, the photosensitive resin layer is exposed patternwise via a pattern mask having a specific pattern;
 in a third step, the photosensitive color resin layer is developed after completion of patternwise exposure, to thereby retain the patterned photosensitive resin layer serving as a pixel layer on the transparent substrate; and
 in a fourth step, the transparent substrate on which a pixel layer is formed is baked (post-baked) after completion of development. A method for producing the color filter of the present invention is not limited to the above-described method so long as the photosensitive coloring composition of the present invention is used.

[0048] Preferred steps for producing a color filter in the present invention will next be described in detail.

[0049] In the first step, the photosensitive coloring composition is applied to a transparent substrate such as a glass substrate through a coating method such as spray-coating, spin-coating, roll-coating, or screen-coating. In order to impart suitable fluidity to such a coating apparatus, the photosensitive coloring composition may contain, in addition to the aforementioned solvents, extender pigments such as barium sulfate, calcium carbonate, and silica; and a small amount of silicon-type or fluorine-type surfactants serving as leveling agents or defoaming agents. The applied photosensitive coloring composition may be dried (pre-baked) typically at 30 to 100°C for 10 to 30 minutes so as to remove solvent by use of a blower-equipped oven or a hot plate. When the temperature is very high or the drying time is very long, partial polymerization or cross-linking may occur and the solubility of unexposed portions to a developer may decrease, which disadvantageously causes poor development.

[0050] In the second step, the photosensitive resin layer obtained in the first step is irradiated with a UV beam via a photomask typically having a color filter pattern. A light source such as an ultra-high-pressure Hg lamp or a metal halide lamp is typically employed. Since UV light passing through the photomask typically has a wavelength of 300 nm or more, three types of rays included in the bright line spectrum of the ultra-high-pressure Hg lamp; i.e., i-ray (365 nm), h-ray (405 nm), and g-ray (436 nm), are used for polymerization or cross-linking. Therefore, sensitivity to the above three spectral lines is important when the photosensitive coloring composition is used in forming a color filter.

[0051] In the third step, the exposed photosensitive resin layer obtained in the second step is developed with a weak alkaline developer; i.e., the unexposed (uncured) portion of the photosensitive resin layer is removed so as to provide a pixel layer.

[0052] In the fourth step, the pixel-layer-formed transparent substrate obtained in the third step is post-baked at 160 to 300°C for about 20 to 60 minutes, to thereby obtain a pixel layer of a first color.

[0053] Beyond the first color, color filters of a variety of colors are produced by repeating the above four steps.

EXAMPLES

[0054] The present invention will next be described by way of examples. In the examples, the term "parts" refers to

"parts by weight," and "%" refers to "wt. %." In addition, the prepared photosensitive coloring composition will be referred to as "resist" for convenience of description. In the examples, a alkali-development-type black resist produced by means of a pigment dispersion method, which is widely employed as a method for producing a color filter for a color liquid crystal display at the present time, will be described. Prior to the examples, an example synthesis of an aqueous alkaline solution-soluble resin will be described.

Production Example of Resin Solution 1

[0055] Cyclohexanone (350 parts), styrene (26 parts), 2-hydroxyethyl acrylate (23 parts), methacrylic acid (35 parts), methyl methacrylate (21 parts), and butyl methacrylate (70 parts) were charged into a 1-liter four-neck flask and heated to 90°C. Independently, cyclohexanone (290 parts), styrene (26 parts), 2-hydroxyethyl acrylate (23 parts), methacrylic acid (35 parts), methyl methacrylate (21 parts), butyl methacrylate (70 parts), and azobisisobutyronitrile (1.75 parts) were mixed to prepare a solution. The thus-prepared solution was added dropwise to the above reaction mixture over a three-hour period. The resultant mixture was further allowed to react for three hours at 90°C. Subsequently, a solution of azobisisobutyronitrile (0.75 parts) in cyclohexanone (10 parts) was further added thereto, and the mixture was allowed to react for one hour, to thereby synthesize a resin solution. A portion of the solution was sampled and heated at 180°C for 20 minutes for drying, to thereby quantitatively determine non-volatile component content. Based on the measurement, cyclohexanone was added to the synthesized resin solution so as to adjust the content of non-volatile component to 20%. Thus, Resin Solution 1 was prepared.

Production Example of Resin Solution 2

[0056] Cyclohexanone (350 parts), styrene (26 parts), 2-hydroxyethyl acrylate (44 parts), acrylic acid (35 parts), and butyl methacrylate (70 parts) were charged into a 1-liter four-neck flask and heated to 90°C. Independently, cyclohexanone (290 parts), styrene (26 parts), 2-hydroxyethyl acrylate (44 parts), acrylic acid (35 parts), butyl methacrylate (70 parts), and azobisisobutyronitrile (1.75 parts) were mixed to prepare a solution. The thus-prepared solution was added dropwise to the above reaction mixture over a three-hour period. The resultant mixture was further allowed to react for three hours at 90°C. Subsequently, a solution of azobisisobutyronitrile (0.75 parts) in cyclohexanone (10 parts) was further added thereto, and the mixture was allowed to react for one hour, to thereby synthesize a resin solution. Thereafter, the temperature in the flask was cooled to 80°C, and a solution of isocyanate ethyl methacrylate (24 parts) and tin octylate (0.11 parts) in cyclohexanone (20 parts) was further added dropwise thereto over 10 minutes, and the mixture was allowed to react for 20 minutes, to thereby synthesize a resin solution. In the same manner as for Resin Solution 1, cyclohexanone was added to the synthesized resin solution, to thereby prepare Resin Solution 2 having a non-volatile component content of 20%.

Examples 1 to 20

[0057] Preparation of a black resist

Resin Solution 1: 55 parts
 Special Black (which is manufactured by Degussa): 5.7 parts
 Dispersing agent: 0.29 parts
 Cyclohexanone: 7.8 parts

[0058] The above components were mixed, and dispersed by use of a paint shaker for 24 hours, to thereby prepare a black dispersion. Next, the following components:

Black dispersion: 54.15 parts
 NK Ester ATMPT (which is manufactured by Shin-Nakamura Chemical Co.,Ltd.): 4.85 parts
 Sensitizer: 0.7-1.8 parts
 Organic boron complex: 0.9 parts
 Thiol compound: 0.9 parts
 Cyclohexanone: 39.0 parts

were sufficiently mixed in a vessel, and the resultant mixture was filtered through a 1.0- μm filter, to thereby prepare a black resist having a non-volatile component content of about 20%.

[0059] Combinations of the sensitizer, organic boron complex, and thiol compound and the proportions of compo-

nents in the sensitizer are shown in Table 1 below.

5

10

15

20

25

30

35

40

45

50

55

Table 1

	Sensitizer (0.7-1.8 parts)	Organic Boron Complex (0.9 parts)	Thiol Compound (0.9 parts)
Example 1	Basic Yellow 21 (0.3 parts) Ethyl Michler's ketone (0.4 parts) HABI (1.0 part)	Tetrabutylammonium butyltriphenoxyborate	2-Mercapto-benzothiazole
Example 2	Basic Yellow 21 (0.3 parts) Ethyl Michler's ketone (0.4 parts) HABI (1.0 part)	Tetrabutylammonium butyltri(4-methylnaphthyl)borate	2-Mercapto-benzothiazole
Example 3	Basic Yellow 21 (0.3 parts) Ethyl Michler's ketone (0.4 parts) HABI (1.0 part)	Tetrabutylammonium methyltri(4-methylnaphthyl)borate	2-Mercapto-benzothiazole
Example 4	Basic Yellow 21 (0.3 parts) Ethyl Michler's ketone (0.4 parts) HABI (1.0 part)	Tetrabutylammonium methyltri(4-methylnaphthyl)borate	Penta-erythritol tetrakis(3-mercaptopropionate)
Example 5	Basic Yellow 21 (0.3 parts) HABI (1.0 part)	Tetrabutylammonium methyltri(4-methylnaphthyl)borate	2-Mercapto-benzothiazole
Example 6	Ethyl Michler's ketone (0.3 parts) HABI (1.0 part)	Tetrabutylammonium methyltri(4-methylnaphthyl)borate	2-Mercapto-benzothiazole
Example 7	Basic Yellow 21 (0.3 parts) Ethyl Michler's ketone (0.4 parts) HABI (1.0 part)	Tetrabutylammonium methyltri(4-methylnaphthyl)borate	2-Mercapto-thiazoline
Example 8	Basic Yellow 13 (0.3 parts) Ethyl Michler's ketone (0.4 parts) HABI (1.0 part)	Tetrabutylammonium methyltri(4-methylnaphthyl)borate	2-Mercapto-benzothiazole
Example 9	Basic Yellow 21 (0.3 parts) Ethyl Michler's ketone (0.3 parts) Benzophenone (0.2 parts) HABI (1.0 part)	Tetrabutylammonium methyltri(4-methylnaphthyl)borate	2-Mercapto-benzothiazole

5
10
15
20
25
30
35
40
45
50
55

Example 10	Basic Yellow 21 (0.3 parts) Ethyl 4-dimethylamino-benzoate (0.3 parts) HABI (1.0 part)	Tetrabutylammonium methyltri(4-methyl-naphthyl)borate	2-Mercapto-benzothiazole
Example 11	Ethyl Michler's ketone (0.3 parts) HABI (1.0 part)	Tetrabutylammonium methyltriphenoxyborate	2-Mercapto-benzothiazole
Example 12	Ethyl Michler's ketone (0.3 parts) HABI (1.0 part)	Tetrabutylammonium methyltrifluorophenylborate	2-Mercapto-benzimidazole
Example 13	Ethyl Michler's ketone (0.3 parts) HABI (1.0 part)	Tetrabutylammonium methyltrifluorophenylborate	Trimethylol-propane tris(3-mercaptopropionate)
Example 14	Ethyl Michler's ketone (0.3 parts) HABI (1.0 part)	Tetrabutylammonium methyltrifluorophenylborate	Penta-erythritol tetrakis(3-mercaptopropionate)
Example 15	Ethyl Michler's ketone (0.3 parts) HABI (1.0 part)	Tetrabutylammonium methyltrifluorophenylborate	Trimethylol-propane tris(3-mercaptopropionate)
Example 16	Ethyl Michler's ketone (0.3 parts) HABI (1.0 part)	Tetrabutylammonium methyltris(4-methyl-naphthyl)borate	Trimethylol-propane tris(3-mercaptopropionate)
Example 17	Basic Yellow 21 (0.3 parts) HABI (1.0 part)	Tetrabutylammonium methyltri(4-methyl-naphthyl)borate	Trimethylol-propane tris(3-mercaptopropionate)
Example 18	Basic Yellow 21 (0.3 parts) Ethyl Michler's ketone (0.3 parts) HABI (1.0 part)	Tetrabutylammonium methyltri(4-methyl-naphthyl)borate	Trimethylol-propane tris(3-mercaptopropionate)
Example 19	Ethyl Michler's ketone (0.3 parts) Benzophenone (0.3 parts) HABI (1.0 part)	Tetrabutylammonium methyltri(4-methyl-naphthyl)borate	Trimethylol-propane tris(3-mercaptopropionate)
Example 20	Ethyl 4-dimethylamino-benzoate (0.3 parts) HABI (1.0 part)	Tetrabutylammonium methyltri(4-methyl-naphthyl)borate	Trimethylol-propane tris(3-mercaptopropionate)

Comparative Example

[0060] The following components:

- 5 The above black dispersion: 54.15 parts
 NK Ester ATMPT (which is manufactured by Shin-Nakamura Chemical
 Co., Ltd.): 4.85 parts
 10 2-(4-Methoxyphenyl)-4,6-bis(trichloromethyl)-s-triazine: 2 parts
 Cyclohexanone: 39.0 parts

10 were sufficiently mixed in a vessel, and the resultant mixture was filtered through a 1.0- μm filter, to thereby prepare a black resist having a non-volatile component content of about 20%, to serve as a comparative example.

15 [0061] In order to measure the spectral sensitivity of the resists obtained, each resist was applied to a glass plate (100 mm x 100 mm) by use of a spin-coater so as to obtain a film having a dry thickness of 1.4 μm , and was dried at 70°C for 20 minutes in a blower-equipped oven. The resist was irradiated with a beam generated from a spectral irradiator (JASCO Model CT-25CP, manufactured by Nihonbunko). An ultra-high-pressure Hg lamp was used as a light source. After completion of exposure, the substrate was immersed in a 1 % aqueous solution of sodium carbonate for approximately 60 seconds for development, washed in a water stream, and heated at 220°C for 30 minutes, to thereby obtain a spectral photograph. In Table 3, the step tablet numbers at which the cured composition remained after development are shown for the cases of exposure to i-ray (365 nm), h-ray (405 nm), and g-ray (436 nm), respectively. The relationship between the step and light intensity is shown in the following Table 2. Briefly, the greater the number of the step, the higher the sensitivity.

Table 2

Step No. (steps)	Light intensity (mJ/cm ²)
13	1.00
12	1.78
11	3.16
10	5.62
9	10.0
8	17.8
7	31.6
6	56.2
5	100
4	178
3	316
2	562
1	1000

45 The results are shown in Table 3.

Table 3

Example No.	i-ray	h-ray	g-ray
Example 1	3	3	3
Example 2	5	4	4
Example 3	6	6	5
Example 4	10	8	7
Example 5	3	6	5

Table 3 (continued)

Example No.	i-ray	h-ray	g-ray
Example 6	2	4	4
Example 7	6	6	5
Example 8	6	5	5
Example 9	7	6	5
Example 10	6	6	5
Example 11	5	4	0
Example 12	5	4	0
Example 13	8	7	1
Example 14	8	7	1
Example 15	9	8	2
Example 16	12	9	2
Example 17	6	9	8
Example 18	12	9	8
Example 19	12	10	2
Example 20	11	9	2
Comparative Example	3	0	0

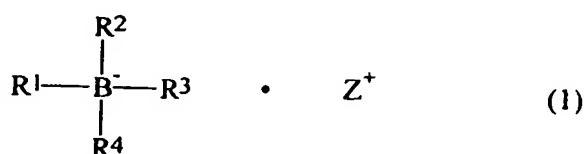
[0062] As is apparent from the results in Table 3, according to the present invention, a photosensitive coloring composition of high sensitivity can be obtained without formation of an oxygen-shielding film. Therefore, steps in a process for producing a color filter can be reduced, and cost reduction is attained due to enhancement of productivity. In addition, a color filter having excellent resistance can be obtained.

[0063] While the invention has been described in detail and with reference to specific embodiments thereof, it will be apparent to one skilled in the art that various changes and modifications can be made therein without departing from the spirit and scope thereof.

Claims

1. A photopolymerization initiator for producing a color filter characterized by comprising:

a sensitizer (A) which absorbs light having a wavelength of 300 to 500 nm;
an organic boron complex (B) represented by formula (1):



wherein each of R¹, R², R³, and R⁴ represents an alkyl group, an aryl group, an aralkyl group, an alkenyl group, a heterocyclic group, or an alicyclic group; and Z⁺ represents an ammonium cation, a sulfonium cation, an oxosulfonium cation, a pyridinium cation, a phosphonium cation, an oxonium cation, or an iodonium cation; and
a thiol-group-containing compound (C).

2. The photopolymerization initiator according to claim 1, wherein the thiol-group-containing compound (C) is at least

one member selected from the group consisting of 2-mercaptopbenzothiazole, trimethylolpropane tris(2-mercaptopacetate), trimethylolpropane tris(3-mercaptopropionate), pentaerythritol tetrakis(2-mercaptopacetate), and pentaerythritol tetrakis(3-mercaptopropionate).

- 5 3. The photopolymerization initiator according to claim 1 or 2, wherein the sensitizer (A) contains a benzophenone compound, a biimidazole compound or a mixture thereof.
- 10 4. A photosensitive coloring composition for producing a color filter, which comprising:
- 15 a transparent polymer (D) soluble in a solvent or an aqueous alkaline solution;
 a compound having an ethylenic unsaturated bond (E);
 a pigment (F); and
 a polymerization initiator according to any one of claims 1 to 3.
- 20 5. A color filter having pixels formed on a transparent substrate by use of the photosensitive coloring composition according to claim 4.
- 25 6. A method for producing a color filter comprising the steps of:
 forming a layer of the photosensitive coloring composition according to claim 4 on a transparent substrate; and
 photocuring an exposed portions of the layer of the photosensitive coloring composition.
- 30 7. The photopolymerization initiator according to claim 1, wherein, based on the compound having an ethylenic unsaturated bond, the sensitizer (A) is in an amount of 1 to 60% by weight, an organic boron complex (B) is in an amount of 1 to 60% by weight, and a thiol-group-containing compound (C) is in an amount of 1 to 60% by weight.
- 35 8. The photopolymerization initiator according to claim 1, wherein the organic boron complex (B) is used in an amount of 5 to 500% by weight, based on the sensitizer (A) and the thiol-group-containing compound (C) is used in an amount of 10 to 500% by weight, based on the organic boron complex (B).
- 40 9. The photopolymerization initiator according to claim 1, wherein the organic boron complex (B) is used from 20 to 300% by weight, based on the sensitizer (A) and the thiol-group-containing compound (C) is used from 50 to 200% by weight, based on the organic boron complex (B).

35

40

45

50

55

(19)



Europäisches Patentamt
European Patent Office
Office européen des brevets



(11)

EP 1 031 579 A3

(12)

EUROPEAN PATENT APPLICATION

(88) Date of publication A3:
06.12.2000 Bulletin 2000/49

(51) Int Cl.7: C08F 2/50, G03F 7/029,
G03F 7/00

(43) Date of publication A2:
30.08.2000 Bulletin 2000/35

(21) Application number: 00103990.8

(22) Date of filing: 25.02.2000

(84) Designated Contracting States:
AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE

Designated Extension States:
AL LT LV MK RO SI

(30) Priority: 26.02.1999 JP 4974299
04.05.1999 US 132449 P

(71) Applicant: SHOWA DENKO KABUSHIKI KAISHA
Minato-ku, Tokyo (JP)

(72) Inventors:

- Katoh, Tsuyoshi, c/o Showa Denko K.K.
Cent.Res.Lab
Kawasaki-shi, Kanagawa-ken (JP)
- Ogata, Tomonari, c/o Showa Denko K.K.
Cent.Res.Lab
Kawasaki-shi, Kanagawa-ken (JP)

(74) Representative: Strehl Schübel-Hopf & Partner
Maximilianstrasse 54
80538 München (DE)

(54) **Photopolymerization initiator for color filter, photosensitive coloring composition, and color filter**

(57) A photopolymerization initiator for producing a color filter including a sensitizer (A) which absorbs light having a wavelength of 300 to 500 nm, an organic boron complex (B), and a thiol-group-containing compound (C); a photosensitive coloring composition including a transparent polymer (D) which is soluble to a solvent or

an aqueous alkaline solution, a compound having ethylenic unsaturated bonds (E), a pigment (F), and the photopolymerization initiator; a color filter having a pixel formed by use of the coloring composition on a transparent substrate; and a method for producing the color filter.

EP 1 031 579 A3



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number

EP 00 10 3990

DOCUMENTS CONSIDERED TO BE RELEVANT									
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.)						
X	EP 0 889 362 A (SHOWA DENKO KK) 7 January 1999 (1999-01-07) * page 8, line 13 - line 18 * * claims *	1-4	C08F2/50 G03F7/029 G03F7/00						
P, X	EP 0 900 800 A (SHOWA DENKO KK) 10 March 1999 (1999-03-10) * claims 1,6-16 *	1-4							
X	EP 0 879 829 A (SHOWA DENKO KK) 25 November 1998 (1998-11-25) * example 3 *	1-3, 7-9							
X	LI ET AL: "4-(4'-dimethylaminostyryl)benzophenone/di phenyliodonium salt - a new kind of electron transfer photosensitization system for radical photopolymerization" CHEMICAL ABSTRACTS + INDEXES, US, AMERICAN CHEMICAL SOCIETY, COLUMBUS, vol. 121, no. 18, 31 October 1994 (1994-10-31), XP002139194 ISSN: 0009-2258 * abstract *	1-3							
X	GB 2 307 474 A (CIBA GEIGY AG) 28 May 1997 (1997-05-28) * page 28, line 19 - line 26 * * claims 1,16,17 *	1-3							
A	DE 44 43 780 A (FUJI PHOTO FILM CO LTD) 14 June 1995 (1995-06-14) * the whole document *	4-6							
A	US 5 622 794 A (SATO MORIMASA ET AL) 22 April 1997 (1997-04-22) * the whole document *	4-6							
<p>The present search report has been drawn up for all claims</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 33%;">Place of search</td> <td style="width: 33%;">Date of completion of the search</td> <td style="width: 34%;">Examiner</td> </tr> <tr> <td>THE HAGUE</td> <td>12 October 2000</td> <td>Pollio, M</td> </tr> </table> <p>CATEGORY OF CITED DOCUMENTS</p> <p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document</p>				Place of search	Date of completion of the search	Examiner	THE HAGUE	12 October 2000	Pollio, M
Place of search	Date of completion of the search	Examiner							
THE HAGUE	12 October 2000	Pollio, M							

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 00 10 3990

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on. The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

12-10-2000

Patent document cited in search report		Publication date		Patent family member(s)	Publication date
EP 0889362	A	07-01-1999	JP US	11072915 A 6033829 A	16-03-1999 07-03-2000
EP 0900800	A	10-03-1999	JP	2000086670 A	28-03-2000
EP 0879829	A	25-11-1998	JP	10316708 A	02-12-1998
GB 2307474	A	28-05-1997	US AT AT AU AU BE BR CA CN DE ES FR IT JP NL NL SG	5952152 A 406775 B 204096 A 717137 B 7179596 A 1010761 A 9605697 A 2191050 A 1158854 A 19648313 A 2126499 A 2741622 A MI962438 A 9188685 A 1004597 C 1004597 A 43445 A	14-09-1999 25-08-2000 15-01-2000 16-03-2000 29-05-1997 05-01-1999 18-08-1998 25-05-1997 10-09-1997 28-05-1997 16-03-1999 30-05-1997 22-05-1998 22-07-1997 07-01-1998 27-05-1997 17-10-1997
DE 4443780	A	14-06-1995	JP US	7159991 A 5622794 A	23-06-1995 22-04-1997
US 5622794	A	22-04-1997	JP DE	7159991 A 4443780 A	23-06-1995 14-06-1995

EPO FORM P0459

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82